

Ophthalmic and ophthalmologic use of a complex
nutritive base in an aqueous medium

5 The present invention relates to the ophthalmic and
ophthalmologic use of a complex nutritive base both in
man and in animals.

10 "Ophthalmic" is to be understood as meaning that a
complex nutritive base as described and defined below
may be utilized in various nontherapeutic applications
in relation to the human or animal eye and, more
specifically, in relation to the external surface or
exterior of the cornea.

15 "Ophthalmologic" is to be understood as meaning that
the same complex nutritive base may be utilized in the
therapeutic or clinical treatment of the human or
animal eye and, more specifically, for application,
e.g. local application, in external contact with the
20 cornea.

More specifically, for the purposes of the uses defined
above, the present invention is concerned with trophic
compositions comprising a complex nutritive base.

25 Complex nutritive base means any composition or
formulation in an aqueous medium which differs from a
cell culture medium in the manner specified below, even
if, like the latter, it generally allows viable in-
30 vitro culturing of an inoculum of certain predetermined
cells for at least 72 h.

Various culture media have actually been described and
are available commercially. Thus, in regard to the in-
35 vitro culturing of keratinocytes, mention may be made
of the following:

- BOYCE ST, HAM RG, Calcium-regulated differentiation
of normal human epidermal keratinocytes in defined

clonal culture and serum-free serial culture; J. Invest. Dermatol. 1983; 81: 335-409;

5 - BOYCE ST, HAM RG, Cultivation, frozen storage, and clonal growth of normal human epidermal keratinocytes in serum-free media; J. Tissue Culture Methods 1985; 9; 83-93;

10 - the commercial medium known as MCDB 153, marketed by companies such as IRVINE SCIENTIFIC and GIBCO-BRL;

- the commercial media known as DMEM (DUBECO Modified Epidermal Medium) and KSFM from GIBCO-BRL, etc.

15 To be effective, culture media of the above kind incorporate cellular growth factors, which may already be present in the composition of the culture medium at the outset or may be produced during culturing, e.g. by a feeder strain of fibroblasts, it being a question of
20 the culturing of keratinocytes.

The use of a complex nutritive base of the kind considered in the case of the present invention does not involve growth factors, such as EGF (Epidermal
25 Growth Factor) for example.

Many of the above-mentioned culture media include biological extracts, e.g. of animal, cellular, or other origin, i.e. ones obtained from a biological starting
30 material. Examples of these biological extracts are any fetal calf serum and any bovine pituitary stalk extract.

By nature, these extracts are of varying, indeed
35 uncertain composition.

A complex nutritive base of the kind considered in the case of the present invention does not include any biological extract as defined above.

In certain cases these same culture media mentioned above include various therapeutic active principles used as medicines, to aid the preservation and/or efficacy of the culture medium.

Examples of such active principles are certain antibiotics, e.g. penicillin and/or streptomycin, alone or in combination, and certain hormones, e.g. cholera toxin and insulin.

A complex nutritive base of the kind considered in the case of the present invention does not include any medicinal active principle.

Complex nutritive bases of this kind, in the sense of ones which may be used, alone or in combination with other constituents, as an active product or as an excipient, have been described in document WO 96/21421.

Complex nutritive bases of this kind comprise, in an aqueous medium, at least a multiplicity of amino acids, some of which are essential amino acids, various vitamins, trace elements, and metallic salts.

In document WO 96/21421, a complex nutritive base of this kind has, for example, the following composition, as shown in Table 1 below:

Table 1

CONSTITUENTS	Concentration in mg/l
Water	q.s.
<u>Amino acids</u>	
L-Alanine	9.2
L-Arginine HCl	421.4
<u>Amino acids (continued)</u>	
L-Asparagine (anhydrous)	14.2
L-Aspartic acid	4.0
L-Cysteine HCl.H ₂ O	42.0
L-Glutamic acid	14.8
L-Glutamine	1754.4
Glycine	7.6
L-Histidine HCl.H ₂ O	50.0
L-Isoleucine	6.0
L-Leucine	131.2
L-Lysine HCl	54.0
L-Methionine	13.5
L-Phenylalanine	10.0
L-Proline	34.6
L-Serine	126.1
L-Threonine	24.0
L-Tryptophan	9.3
L-Tyrosine 2 Na 2H ₂ O	11.7
L-Valine	70.3
<u>Vitamins</u>	
d-Biotin	0.02
Folic acid	0.80
Nicotinamide	0.04
D-Ca pantothenate	0.30
Pyridoxine HCl	0.06
Riboflavin	0.04
Thiamine HCl	0.30
i-Inositol	18.0
Sodium pyruvate	55.0
Thymidine	0.73

CONSTITUENTS	Concentration in mg/l
Adenine (HCl)	24.0
DL-Lipoic acid	0.20
<u>Inorganic constituents</u>	
Sodium chloride	6800.0
KCl	112.0
Na ₂ HPO ₄	284.0
CuSO ₄ .5H ₂ O	0.003
Sodium acetate	300.0 (anhydrous)
D-Glucose	1080.0
HEPES (piperazine)	6600.0
Phosphorylethanolamine	0.06768
Ethanolamine	0.04684
Sodium sulfate	3.4
Sodium bicarbonate	1160.0
FeSO ₄ .7H ₂ O	1.39
MgCl ₂ .6H ₂ O	120.0
CaCl ₂ .2H ₂ O	13.0-22.05
ZnSO ₄ .7H ₂ O	0.144
(NH ₄) ₆ Mo ₇ O ₂₄ .4H ₂ O	0.00120
Na ₂ SiO ₃ .5H ₂ O	0.142
<u>Inorganic constituents</u> (continued)	
MnCl ₂ .4H ₂ O	0.00002
SnCl ₂ .2H ₂ O	0.00011
NH ₄ VO ₃	0.00057

Document WO 02/087326 describes and proposes an ophthalmic and ophthalmologic composition comprising a vitamin derivative as a nonionic surfactant and a cationic agent or preservative, e.g. polyhexamethylene biguanide, and a disinfectant, e.g. hydrogen peroxide. This composition essentially cleans and disinfects, but does not regenerate the cornea.

10 Document US 5 654 266 describes and proposes an essentially buffered saline composition for preserving

and washing tissues prior to transplantation, e.g. corneas to be used for grafts. The proposed composition comprises mainly β -hydroxybutyrate, reducing the production and accumulation of lactic acid in the stored tissues.

Document EP 0 517 970 describes and proposes a solution for intraocular irrigation of the anterior and posterior chambers of the eye, for ocular surgery purposes, which repairs solely the corneal endothelium, i.e. the anterior chamber of the eye.

Document JP 02 083318 describes a composition which cleans the eye, comprising quaternary ammonium salts, chondroitin-sodium sulfate, glycyrrhetic acid, and alcohols, including propylene glycol.

Document WO 026495 describes a composition for the care, and particularly the impregnation, of contact lenses, which contains vitamins, the aim being that the said composition be released from the impregnated contact lens on contact with the cornea.

The subject of the present invention is the ophthalmic or ophthalmologic use of a complex nutritive base as defined above, e.g. via a trophic composition comprising said base, notably to improve viability, e.g. under conditions of attack or stress, and to maintain the integrity and equilibrium of the epithelial cells of the cornea of the eye, in man or animals.

More specifically, the subject of the present invention is a use of the above kind to improve the viability, growth, and differentiation of corneal epithelial cells, in man or animals.

In the present invention, the trophic composition in an aqueous medium which may be utilized comprises:

- a complex nutritive base comprising at least a multiplicity of amino acids, vitamins, trace elements, and metallic salts, the said base being free of any cellular growth factor, any biological extract of animal or cellular origin, or any therapeutic active principle,

- an inhibitor of collagenases of the human or animal corneal epithelium, and

- a promoter of neocollagen synthesis.

Preferably the trophic composition is formulated with the complex nutritive base to establish a pH between 7.3 and 7.5 and an osmolarity between 300 and 350 mOsm.

Various collagenase inhibitors are already known, functionally and/or chemically; reference may be made to the following documents in particular:

- Berman; Int. Ophtalmol. Clin. 1975 Winter; 15(4): 49-66
- Haffner JC, Fecteau KA, Eilet H, Vet. Ophtalmol. 2003 Mar; 6(1): 67-72.

Various promoters of neocollagen synthesis are already known; reference may be made to the following documents in particular:

- Bellon G, Monboisse JC, Fandoux A, Borel JP, Biochim. Biophys. Acta. 1987, Aug 19, 930 (1): 39-47
- Kefalides NA, Cameron JD, Tomicheck EA, Yanoff M, J Biol Chem. 1976, Feb 10; 251(3): 730-3.

According to the invention, the trophic composition benefits from the following characteristics, the latter being considered alone or in combination:

- the inhibitor of collagenases is chosen from the group comprising cysteine, N-acetylcysteine, and EDTA calcium salt;
- the inhibitor of collagenases is N-acetylcysteine;
- 5 - the inhibitor of collagenases represents at most 5% and preferably between 0.05 and 0.5% by weight of the said composition;
- the promoter of neocollagen synthesis is proline or hydroxyproline;
- 10 - the promoter of neocollagen synthesis represents at most 0.5% and preferably 0.004% by weight of the said composition;
- the trophic composition comprises hyaluronic acid and/or a salt of hyaluronic acid in a total proportion
- 15 by weight of the composition of at most 0.1% and preferably 0.07%;
- the trophic composition includes a preservative in a proportion by weight of the composition of at most 0.0001%;
- 20 - the preservative is polyhexanide or polyhexamethylene biguanide (PHMB);
- the trophic composition corresponds to the formula described by Table 2 in the description.
- 25 Compared to the composition as per Table 1, the composition as per Table 2 differs from the latter in the following respects:
 - it does not include vitamin B12, putrescine 2HCl, or
 - 30 Na_2SiO_3
 - it includes both a collagenase inhibitor and a promoter of neocollagen synthesis
 - its pH and osmolarity characteristics allow oxygenation of the corneal epithelium and tear lysozyme
 - 35 activity.

The invention also relates to the use of a trophic composition as described above as an ophthalmologic and contactologic medicine in man or animals, by

application, e.g. local application, in external contact with the cornea.

5 It also relates to such a medicine in liquid form or in dry form, that is to say for reconstitution with an aqueous medium.

10 In one variant, such a medicine is in liquid form, e.g. in the form of drops, or regenerating tears, or eyewash, or solution.

15 Thus, as an example, the present invention proposes an eyewash comprising a trophic composition as defined above, which can heal the cornea or prevent conjunctival adhesions.

Such an eyewash has the following indications:

- 20 - corneal ulcers of traumatic or other origin, corneal burns;
- prevention of conjunctival adhesions and corneoconjunctival adhesions (symblepharon);
- conjunctival and corneal xerosis.

25 The invention also relates to an ophthalmic solution, e.g. comfort solution, for local application in external contact with the cornea in man or animals, comprising for example a trophic composition as defined above.

30 Thus the present invention proposes, for example, "regenerating tears" comprising a trophic composition as defined above for the hydration of contact lenses and regeneration of the cornea in man or animals.

35 "Regenerating tears" of this kind also lubricate the cornea at the same time. In the final analysis, they prevent or limit premature erosion of the cornea as a result of the action of contact lenses, hard or semi-

rigid ones in particular. These "regenerating tears" both improve visual comfort and increase the length of time for which contact lenses can be worn.

5 Thus the present invention proposes, for example, comfort drops for the following uses:

- ocular dryness of iatrogenic origin (associated, in particular, with the use of topical or systemic
10 retinoids, antihistamines, antiparkinsonian drugs, beta blockers, spasmolytics, anxiolytics, neuroleptics, antidepressants, and bronchodilators, etc.), producing discomfort, particularly in patients who wear contact lenses;

15

- ocular dryness in the elderly.

The invention also relates to the use of a complex nutritive base as defined above for the treatment, e.g.
20 the preservation, of items or prostheses, such as contact lenses, which are designed to come into external contact with the cornea of the eye in man or in animals.

25 It relates, in this regard, to a solution for the preservation, or storage, or transport, or use (e.g. surgical use) of items or prostheses, such as contact lenses, which are designed to come into external contact with the cornea of the eye in man or in
30 animals.

Thanks to a complex nutritive base as defined above, and, in particular, to a trophic composition in accordance with the present invention, and as
35 demonstrated by the studies described below, the primary intrinsic properties of the corneal epithelium are preserved for a long time and, at the same time, its resistance to attack is increased and, where

applicable, its return to a state of equilibrium promoted.

The present invention is now described and exemplified
 5 on the basis of a trophic composition in an aqueous medium with the formula shown in Table 2.

Table 2

Line No.	International Nomenclature Cosmetic Ingredient name (INCI name)	Concentration in mg/l	Line No.	International Nomenclature Cosmetic Ingredient name (INCI name)	Concentration in mg/l
1	WATER	q.s.	14	SODIUM PYRUVATE	55
2	SODIUM CHLORIDE	6800	15	LYSINE HCL	54
3	GLUTAMINE	1754.4	16	HISTIDINE HCL	50
4	SODIUM BICARBONATE	1160	17	CYSTEINE HCL	42
5	GLUCOSE	1080	18	ADENINE	24
6	ARGININE HCL	421.4	19	THREONINE	24
7	SODIUM ACETATE	300	20	Ca CHLORIDE	20.05
8	DISODIUM PHOSPHATE	284	21	INOSITOL	18
9	LEUCINE	131.2	22	GLUTAMIC ACID	14.8
10	SERINE	126.1	23	ASPARAGINE	14.2
11	Mg CHLORIDE	120.0	24	METHIONINE	13.5
12	K CHLORIDE	112	25	TYROSINE	11.7
13	VALINE	70.3	26	PHENYLALANINE	10.0
			27	TRYPTOPHAN	9.3

Line No.	International Nomenclature Cosmetic Ingredient name (INCI name)	Concentration in mg/l
28	ALANINE	9.2
29	GLYCINE	7.6
30	ISOLEUCINE	6.0
31	ASPARTIC ACID	4.0
32	SODIUM SULFATE	3.4
33	FERROUS SULFATE	0.003
34	FOLIC ACID	0.8
35	THYMIDINE	0.73
36	CYANOCOBALAMIN	0.41
37	CALCIUM PANTOTHENATE	0.3
38	THIAMINE HCL	0.3
39	THIOCTIC ACID	0.3
40	ZINC SULFATE	0.144
41	SODIUM SILICATE	0.142
42	PYRODIXINE HCL	0.06
43	NIACINAMIDE	0.04
44	RIBOFLAVIN	0.3
45	BIOTIN	0.02
46	COPPER SULFATE	0.003
47	AMMONIUM MOLYBDATE	0.00120
48	AMMONIUM VANADATE	0.003
49	Mn CHLORIDE	0.00002
50	SODIUM HYALURONATE	70
51	POLYHEXANIDE or POLYHEXAMETHYLENE BIGUANIDE	0.1
52	N-ACETYLCYSTEINE	500
53	HYDROXYPROLINE or PROLINE	35

In line with the definition of the trophic composition in accordance with the present invention, the ingredients of the complex nutritive base in an aqueous medium are all those numbered 1-49, whilst ingredients 5 50-53 are considered not to form part of the complex nutritive base, and serve to make it suitable for external contact with the corneal epithelium of the eye.

10 Ingredients 50 and 51 are optional, depending on the role or function sought for the trophic composition.

Moreover, the choice and proportions of the various ingredients are defined so as to establish, in the 15 final trophic composition:

- a pH between 7.3 and 7.5 and preferably between 7.42 and 7.45 (a pH which promotes oxygenation of the corneal epithelium and tear lysozyme activity);
- 20 - an osmolarity between 300 and 350 mOsm, 340 mOsm for example.

Study 1

25 This evaluates the biocompatibility of the trophic composition defined above in a reconstituted human corneal epithelium model.

30 Starting from the formula shown in Table 2, four variants are defined and formulated, namely:

- EVE 1: without sodium hyaluronate (ingredient 50) and without preservative (ingredient 51).
- EVE 2: without sodium hyaluronate (ingredient 50) but with preservative (ingredient 51).
- EVE 3: with sodium hyaluronate (ingredient 50) but without preservative (ingredient 51).
- EVE 4: identical to the formula shown in Table 2,

i.e. with sodium hyaluronate (ingredient 50)
and with preservative (ingredient 51).

Minor adjustments are made to the other ingredients in order to keep the pH and osmolarity within the values stated above.

5

The cytocompatibility of formulas EVE 1 to EVE 4 is evaluated by topical application using an in-vitro reconstructed human corneal epithelium model of the kind available from SKINETHIC LABORATORIES, 45 Rue
10 Saint Philippe, 06000 NICE, France.

Roger W. BEUERMAN and Lia PEDROSA have described in detail the ultrastructure of the human cornea in an article entitled "Ultra-structure of the human cornea"
15 in the journal "Microscopy Research and Technique", 33: 320-335 (1996), to which reference should be made as required.

Starting from the natural structure, O. DOUCET et al.
20 have proposed and described a reconstructed, three-dimensional, human corneal epithelium, for the performance of various in-vitro toxicity studies in particular; cf. the article entitled "A new in vitro human epithelium model for assessing the eye irritation
25 potential of formulated cosmetic products" published in "In vitro and Molecular Toxicology", 11, 4: 273-283 (1998).

This reconstructed corneal epithelium, or model, is
30 obtained by culturing corneal epithelial cells, such as the HCE line available from and marketed by LSU Eye Center, Louisiana State University School of Medicine, New Orleans, Louisiana, 70112 USA. The epithelial cells are cultured at the air/liquid interface in a defined
35 medium and they form a corneal epithelial tissue, without a stratum corneum, which is close to the corneal epithelium in vivo.

- The study is carried out in reconstructed corneal epithelia of this kind as obtained on the 5th day of culturing (size: 0.5 cm²), the quality control of the various epithelia at the various stages of culturing being undertaken by the aforementioned SKINETHIC LABORATORIES, in accordance with the last cited publication.
- 10 The test principle consists in applying the trophic composition in direct contact with the reconstructed corneal epithelium for 15 minutes, three times a day, for a period of 7 days. An untreated control corneal epithelium is run in parallel. Each condition (except
- 15 for the untreated control) is carried out in duplicate. On the 2nd, 4th, and 7th days of the study, cell viability (MTT test) is evaluated for each of the conditions.
- 20 In practice, 30 µl of the trophic compositions to be tested is placed directly on the surface of the corneal equivalents 3 times a day (at 09:00, 13:00, and 17:00) for 15 minutes in each case. After this contact period, excess composition is removed and the epithelia are
- 25 incubated at 37°C, in a humid atmosphere, 5% CO₂. This operation is repeated every day for 7 days. On the 2nd, 4th, and 7th days of the study, samples of the reconstructed epithelia are collected for the cell viability test.
- 30 On the 2nd, 4th, and 7th days of the study, the epithelia samples are rinsed with PBS and placed in 300 µl of MTT (0.5 mg/ml). After 3 hours' incubation at 37°C, 5% CO₂, the epithelia are transferred into 1.5 ml isopropanol.
- 35 The MTT cleavage product is extracted with gentle stirring over a period of 2 h at room temperature. The optical density at 570 nm is then measured in 200 µl of extraction solution. The results are expressed as

percentage viability compared to the (untreated) control:

$$\% \text{ viability} = [\text{OD}_{(570 \text{ nm test product})} / \text{OD}_{(570 \text{ nm control})}] \times 100$$

5

Under the experimental conditions defined, the viability of the reconstructed human corneal epithelia in vitro is fully maintained after topical application of each of the aforementioned trophic compositions. The presence of the preservative and of hyaluronic acid at the specified concentrations of use does not have any effect on the cellular viability of the epithelia, which is maintained.

15 Study 2

A reconstituted human corneal epithelium is cultured in the trophic compositions defined above and the results obtained are compared with those obtained from culturing in a buffered saline solution.

25

The chosen corneal epithelium model is identical to the one described and defined in Study 1 and is cultured for 72 h in the trophic composition being investigated.

Just one trophic composition is used for this study, namely the one specified in Table 2 and referred to as EVE 4 in Study 1.

30 As soon as the epithelia are received, the SKINETHIC nutritive transport medium is removed, and replaced, in the case of one of the epithelia, with 1 ml of the trophic composition being investigated. A second epithelium is placed in a buffered saline solution
35 (Phosphate Buffered Saline, PBS, Ca^{2+} - and Mg^{2+} -enriched) for comparison. The cytocompatibility of the trophic composition being investigated has previously been verified (no cytotoxic effect, cf. preceding study, Study 1). The epithelia are incubated at 37°C, in a

humid atmosphere, 5% CO₂. The media are replaced every day. After 3 days of culturing, samples of the 2 reconstructed epithelia are collected for histological analysis. They are fixed with a 10% formaldehyde solution and then embedded in paraffin. The histology of the hematoxylin/eosin-stained sections is evaluated under an optical microscope.

The histopathological interpretation of the sections includes the thickness and regularity of the epithelial tissue and also the morphology, cohesion, and proliferative capacity of the cells.

As regards the epithelium cultured in the trophic composition for 72 h, analysis of the sample shows an epithelium implanted on the membranal support. It comprises 6 to 7 layers of closely packed cells (the connecting bridges are visible). There is no sign of necrosis or of keratinization of the superficial layer.

In the case of the epithelium cultured in a buffered saline solution for 72 h, analysis of the sample shows the near-total disappearance of the epithelium; only the support is visible. There is no identifiable cell layer.

Under the experimental conditions defined for this study, a reconstructed human corneal epithelium cultured for 72 h in a trophic composition in accordance with the present invention survives and exhibits cellular cohesion. In contrast, in the similar experiment carried out with a buffered saline solution, there is complete degeneration of the epithelium, which is no longer identifiable.

Study 3

The healing properties of trophic compositions in accordance with the invention are evaluated in a reconstituted human cornea model.

The four variants of the trophic composition investigated are identical to those defined and described in Study 1.

The reconstructed corneal epithelium model is identical to that described and defined in Studies 1 and 2.

The healing properties of the trophic compositions investigated are evaluated after topical application in the aforementioned model, in which a mechanical lesion is created beforehand.

The lesions in the surface of the reconstructed corneal epithelia are created with the tip of a scalpel, each of the epithelia being divided along its diameter.

The test principle consists in applying the trophic composition in direct contact with the reconstructed corneal epithelium for 15 mins, three times a day, every day for a period of 9 days, starting immediately after the creation of the lesions. A control in buffered saline solution (PBS) is run in parallel. On the 2nd, 4th, 7th, and 9th days of the study, the histology of the reconstructed tissue is evaluated for each of the conditions. It is compared with the histology of a control, comparison condition corresponding to an untreated, undamaged epithelium.

In practice, 30 µl of the trophic compositions to be tested is placed directly on the surface of the corneal epithelia 3 times a day (at 09:00, 13:00, and 17:00) for 15 mins in each case. After this contact period, excess composition is removed and the epithelia are

incubated at 37°C, in a humid atmosphere, 5% CO₂. This operation is repeated every day for 9 days. On the 2nd, 4th, 7th, and 9th days of the study, samples of the reconstructed epithelia are collected for histological analysis.

On the 2nd, 4th, 7th, and 9th days of the study, the epithelia are fixed with a 10% formaldehyde solution and then embedded in paraffin. The histology of the haematoxylin/eosin-stained sections is evaluated under an optical microscope.

The healing properties of the trophic compositions are evaluated by histological analysis of the sections and observation of epithelial repair at the site of the lesion created by the scalpel. The site of the lesions is marked with India ink on the surface of the corneal epithelia before histological section (transverse section of the epithelium).

The histopathological interpretation of the sections includes the thickness and regularity of the epithelial tissue and also the morphology, cohesion, and proliferative capacity of the cells after the creation of the lesion.

Combined photographs establish, for each condition, the transverse sections of the epithelia performed at the start of treatment (2nd day after the creation of the lesion) and on the last day of treatment (9th day).

In the case of the (untreated, undamaged) Control, a vital epithelium consisting of 6 to 7 layers of closely packed, regularly shaped cells (with appearance of pavement cells) is observed on the 2nd day of culturing. On the 9th day of treatment there is observable thickening of the epithelium, which is made up of around ten layers of cells, which are still closely

packed and regular. A keratinized layer is seen at the surface.

5 In the case of the treatment with EVE 4 (trophic composition in accordance with Table 2), the photograph taken on the 2nd day shows the lesion created in the epithelium exposing the membranal support. The epithelium is vital on both sides of the wound. There is no keratinization. On the 9th day of treatment a
10 continuous, repaired epithelium made up of several layers of non-necrosed, relatively regular cells (which look more rounded than in the case of the Control) is observable at the site marked with India ink. An area of keratinization is seen in the superficial layers.

15 In the case of the treatment with a buffered saline solution (PBS), the damaged area, marked by a considerable loss of epithelial substance, is observable on the 2nd day. On the 9th day, a repaired
20 epithelium made up of only 5 to 6 layers of cells is observable at the site marked, the cells having a rounded and in some instances a turgescient appearance, with vacuoles.

25 In the case of the treatment with compositions EVE 1, EVE 2, and EVE 3, a lesion marked out by a complete loss of substance is observable in the samples collected on the 2nd day of the study. On both sides, the epithelium has many layers and is vital and not
30 keratinized.

On the 9th day, in the case of the EVE 1 formulation, a continuous reconstituted epithelium made up of rounded, fairly regular cells with a number of vacuoles is
35 observable. In the case of the EVE 2 and EVE 3 formulations, the epithelia look identical with closely packed, morphologically regular cells.

However, the epithelia seem to be thinner than those treated with the EVE 4 formulation.

5 Under the experimental conditions defined, repair of the previously damaged human corneal epithelia is observable at the end of the treatments in each of the conditions tested.

10 The EVE 4 formulation (containing hyaluronic acid and preservative) gives the best results. The repaired epithelium is continuous, vital, has many layers, and is made up of closely packed, regular cells.

Study 4

15

The healing properties of trophic compositions in accordance with the present invention are evaluated in comparison with a buffered saline solution.

20 The trophic composition investigated is identical to the one defined in Table 2.

The reconstructed corneal epithelium model is identical to that used in Studies 1 to 3.

25

The healing properties of the trophic compositions are evaluated by topical application in an in-vitro reconstructed human corneal epithelium model, in which a mechanical lesion is created beforehand.

30

The lesions in the surface of the reconstructed corneal epithelia are created with the tip of a scalpel, each of the epithelia being divided along its diameter.

35 The test principle consists in applying the trophic composition in direct contact with the reconstructed corneal epithelium for 15 mins, three times a day, every day for a period of 7 days, starting immediately after the creation of the lesions. A control in

buffered saline solution (Ca^{2+} - and Mg^{2+} -enriched FBS) is run in parallel. On the 2nd, 4th, and 7th days of the study, the histology of the reconstructed tissue is evaluated for each of the conditions. It is compared
5 with the histology of a control, comparison condition corresponding to an untreated, undamaged epithelium.

In practice, 30 μl of the compositions to be tested is placed directly on the surface of the corneal
10 equivalents 3 times a day (at 09:00, 13:00, and 17:00) for 15 mins in each case. After this contact period, excess composition is removed and the epithelia are incubated at 37°C, in a humid atmosphere, 5% CO_2 . This operation is repeated every day for 7 days. On the 2nd,
15 4th, and 7th days of the study, samples of the reconstructed epithelia are collected for histological analysis.

On the 2nd, 4th, and 7th days of the study, the epithelia
20 are fixed with a 10% formaldehyde solution and then embedded in paraffin. The histology of the hematoxylin/eosin-stained sections is evaluated under an optical microscope.

25 The healing properties of the trophic compositions are evaluated by histological analysis of the sections and observation of epithelial repair at the site of the lesion created by the scalpel. The site of the lesions is marked with India ink on the surface of the corneal
30 epithelia before histological section (transverse section of the epithelium).

The histopathological interpretation of the sections includes the thickness and regularity of the epithelial
35 tissue and also the morphology, cohesion, and proliferative capacity of the cells after the creation of the lesion.

Photographs establish, for each condition, the transverse sections of the epithelia performed at the various stages of treatment.

- 5 In the case of the (untreated, undamaged) Control, a vital epithelium consisting of 6 to 7 layers of closely packed, regularly shaped cells (with appearance of pavement cells) is observed on the 2nd day of culturing. On the 7th day of treatment there is observable
- 10 thickening of the epithelium, which is made up of around ten layers of cells, which are still closely packed and regular. A keratinized layer is seen at the surface.
- 15 In the case of the treatment with the trophic composition shown in Table 2, the photograph taken on the 2nd day shows the lesion created in the epithelium exposing the membranal support. The epithelium is vital on both sides of the wound. There is no keratinization.
- 20 On the 4th day, cellular recolonization of the support is observed. The epithelium is made up of 3 to 4 layers of cells, some of which are in the process of division. On the 7th day of treatment a continuous, repaired epithelium made up of several layers of non-necrosed,
- 25 relatively regular cells (which look more rounded than in the case of the Control) is observable at the site marked with India ink. An area of keratinization is seen in the superficial layers.
- 30 In the case of the treatment with a buffered saline solution (PBS), the damaged area, marked by a considerable loss of epithelial substance, is observable on the 2nd day. On the 4th day the epithelium re-forms, consisting of only one to two layers of
- 35 rounded cells. On the 7th day, the epithelium is repaired and is made up of 5 to 6 layers of cells, these having a rounded and in some instances a turgescent appearance, with vacuoles.

Under the experimental conditions defined, the trophic composition allows the reconstitution of a previously damaged human corneal epithelium. The repaired epithelium is continuous, vital, has many layers, and
5 is made up of closely packed, regular cells.

The application of a buffered saline solution likewise allows repair of the epithelium. However, the latter is made up of only 3 to 4 layers of cells, demonstrating
10 that the regeneration is inferior to that observed with the trophic composition. Moreover, the cells are rounded, with a number of vacuoles.